

Deep marine subsurface archaeal communities of the Peru Trench: Sulfate-reducers, Methanogens, and Unknowns

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Community composition of methanogens and sulfate-reducers was studied in cold, organic-rich sediments of the Peru Trench, collected by the Ocean Drilling Program in 2002 (Leg 201, site 1230). We used PCR assays targeting functional genes of methanogens (*mcrA*) and sulfate reducers (*dsrAB*) throughout the sediment column. Methanogen-specific DNA was successfully amplified from a sample at 44.3 mbsf that coincided with a local acetate peak. Sequences fell into a novel cluster, with members of the obligately acetoclastic *Methanosaeta* group as closest cultured relatives. Sequences of sulfate-reducing prokaryotes were obtained from surface sediments and shared highest similarity with cultured members of the archaeal order *Archaeoglobales*.

We searched for unequivocal evidence of metabolically active Archaea by cloning reverse-transcribed 16S rRNA sequences that had been amplified with general archaeal 16S primers, and 16S primers specific to suspected anaerobic methanotrophs (ANME-2). Preliminary BLAST searches of general archaeal sequences from various depths (0-124 mbsf) revealed active populations of Deep-Sea Archaeal Group (DSAG) and Marine Benthic Group-D (MBG-D). None of the detected archaeal groups had closely related cultured representatives. Suspected anaerobic methanotrophic sequences were amplified successfully from several cores in the methane-sulfate transition zone of Site 1230.

The dominance of uncultured archaeal lineages (DSAG, MBG-D) in the Site 1230 clone libraries contrasts with relative scarcity of recognizable methanogens and sulfate-reducers. These uncultured archaeal subsurface lineages do not share the classical key genes of methanogenic and sulfate-reducing pathways (*mcrA* and *dsrAB*), and represent novel deep subsurface organisms with unknown physiology. Our results substantiate the existence of a systematically distinct archaeal community in the deep marine subsurface.